

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (currently amended): An apparatus for detecting dislodgement of a needle inserted into a patient comprising:
a sensor capable of detecting wetness due to blood wherein the sensor includes a capacitive sensor; and
a barrier pad capable of absorbing blood lost from the patient due to the dislodgment of the needle; and
a sensor holder adapted to secure the sensor ~~in juxtaposition and the barrier pad adjacent~~
~~to the needle such that the sensor detects wetness due to blood loss from the~~
~~patient upon dislodgement of the needle, wherein the sensor holder secures the~~
~~barrier pad between the sensor and the needle such that the sensor does not~~
~~contact blood upon detection thereof.~~

Claims 2-4 (canceled).

5. (previously presented): The apparatus of Claim 1 wherein the capacitive sensor includes one or more electrodes.

6. (currently amended): The apparatus of Claim 5 wherein the capacitive sensor is located within the sensor holder such that the sensor ~~does not contact blood upon detection~~
~~thereof detects wetness due to blood loss in the barrier pad.~~
7. (original): The apparatus of Claim 1 wherein the sensor produces a signal upon detection of blood loss.
8. (original): The apparatus of Claim 7 further comprising a control device adapted to receive the signal for monitoring and controlling blood loss due to the dislodgement of the needle during hemodialysis.
9. (original): The apparatus of Claim 8 wherein the control device is attached to the patient.
10. (original): The apparatus of Claim 1 wherein the needle comprises a venous needle.
11. (currently amended): An apparatus for detecting needle dislodgement during hemodialysis comprising:
a sensor holder having a cavity; ~~and~~
a barrier pad capable of absorbing blood lost from the patient due to the dislodgment of the needle; and
a capacitive sensor comprising an electrode enclosed within the cavity of the sensor holder such that the capacitive sensor is capable of detecting wetness absorbed within the barrier pad from blood due to needle dislodgement during hemodialysis

wherein such that the capacitive sensor does not contact blood upon detection thereof.

12. (original): The apparatus of claim 11 wherein the electrode comprises a single plate electrode.
13. (original): The apparatus of claim 11 further comprising a sterile pad such that the capacitive sensor detects wetness due to blood loss into the sterile pad overlying a vascular access region of a venous needle.
14. (currently amended): The apparatus of claim 13 wherein the sensor holder comprises a flexible material that adaptably conforms to the vascular access region such that the capacitive sensor is capable of detecting blood loss due to needle dislodgement.

Claims 15-16 (canceled).

17. (currently amended): An apparatus for controlling blood loss from a patient during hemodialysis comprising:

a barrier pad capable of absorbing blood lost from the patient due to the dislodgment of the needle; and

a sensor capable of detecting wetness due to blood and a sensor holder adapted to secure the sensor and the barrier pad adjacent to the patient such that the sensor produces a signal indicative of wetness detected within the barrier due to blood loss from the patient upon dislodgement of a venous needle inserted into the patient wherein

the sensor includes a capacitive sensor and the capacitive sensor does not contact blood upon detection thereof; and

a controller capable of processing the signal to prevent blood flow through the venous needle such that blood loss from the patient due to dislodgement of the venous needle is minimized.

18. (original): The apparatus of Claim 17 wherein the sensor holder comprises a pad configuration overlying an access region of the venous needle.
19. (original): The apparatus of Claim 17 further comprising a sterile pad overlying an access region of the venous needle such that the sensor detects wetness in the sterile pad due to blood loss from the patient upon venous needle dislodgement.
20. (original): The apparatus of Claim 19 wherein the sensor contacts the sterile pad to detect wetness therein.
21. (original): The apparatus of Claim 19 wherein the sensor is located inside of the sensor holder such that the sensor does not contact the sterile pad upon detecting wetness therein.

Claim 22 (canceled).

23. (original): The apparatus of Claim 17 wherein the controller is in communication with a hemodialysis machine via an electrical communication cable or a cordless interface to minimize blood loss due to venous needle dislodgement.
24. (original): The apparatus of Claim 23 wherein the controller is adapted to monitor one or more hemodialysis treatment parameters including wetness due to blood loss, change in blood flow and detection of arterial air bubbles during hemodialysis.
25. (original): The apparatus of Claim 24 wherein the controller is attached to the patient for electrical connection to the sensor.
26. (original): The apparatus of Claim 24 wherein the controller comprises a display for monitoring each of the parameters.
27. (currently amended): A method of detecting needle dislodgement comprising the steps of:
providing a barrier pad capable of absorbing blood lost from the patient due to the dislodgment of the needle; and
providing a sensor capable of detecting wetness due to blood wherein the sensor includes a capacitive sensor that does not directly contact blood upon detection thereof;
inserting a needle into a patient; and
securing the sensor and the barrier pad to the patient such that the sensor detects blood absorbed within the barrier pad on the patient upon dislodgement of the needle.

Claim 28 (canceled).

29. (original): The method of Claim 27 wherein the needle comprises a venous needle inserted into the patient for hemodialysis.
30. (currently amended): A method of controlling blood loss from a patient due to needle dislodgement comprising the steps of:
providing a sensor capable of detecting wetness due to blood wherein the sensor includes a capacitive sensor;
providing a barrier pad capable of absorbing blood lost from the patient due to the dislodgment of the needle; and
inserting a needle into the patient;
securing the sensor and the barrier pad adjacent to the patient such that the sensor produces a signal indicative of wetness within the barrier pad due to blood loss from the patient upon dislodgement of the needle and does not directly contact blood upon detection thereof; and
processing the signal to prevent blood flow through the venous needle such that blood loss from the patient due to needle dislodgement is minimized.

Claim 31 (canceled).

32. (original): The method of Claim 30 wherein the needle comprises a venous needle inserted into the patient for hemodialysis.
33. (original): The method of Claim 32 wherein the signal is processed for communicating with a hemodialysis machine to minimize blood loss to the patient due to needle dislodgement.
34. (original): The method of Claim 33 wherein the signal is processed to shut-off a blood pump of the hemodialysis machine.
35. (original): The method of Claim 33 wherein the signal is processed to activate a venous line clamp for preventing blood flow via the venous needle.
36. (currently amended): A method of providing dialysis to a patient comprising the steps of:
providing a barrier pad;
providing a sensor capable of detecting wetness due to blood wherein the sensor includes a capacitive sensor that does not contact blood upon detection thereof;
inserting a venous needle into the patient;
securing the sensor and the barrier pad in juxtaposition to the venous needle;
passing blood through the venous needle via a hemodialysis machine; and
detecting wetness within the barrier pad indicative of blood loss from the patient upon dislodgement of the venous needle such that the sensor does not directly contact blood.

37. (currently amended): The method of Claim 36~~Claim 38~~ wherein blood flow through the venous needle is stopped upon detecting dislodgement of the venous needle such that blood loss from the patient is minimized.

Claim 38 (canceled).